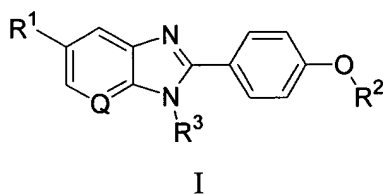


Amendments to the Claims

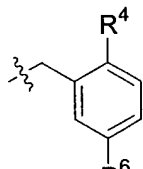
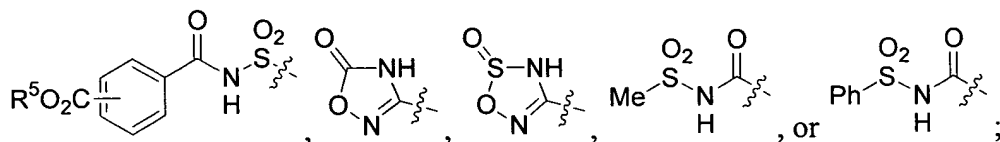
1. (Currently Amended) A compound of Formula I



wherein:

Q is CH or N;

R<sup>1</sup> is tetrazolyl, MeCONHSO<sub>2</sub><sup>-</sup>, PhCONHSO<sub>2</sub><sup>-</sup>, R<sup>5</sup>O<sub>2</sub>C(CH<sub>2</sub>)<sub>0-3</sub>CONHSO<sub>2</sub><sup>-</sup>,

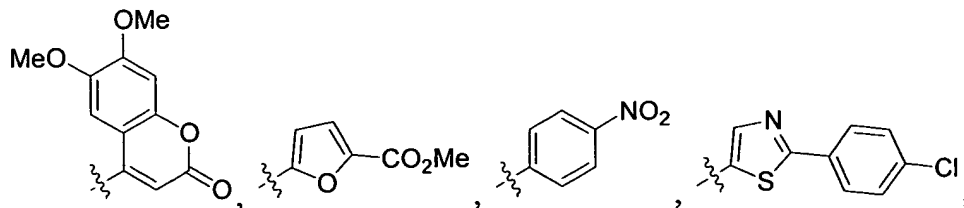
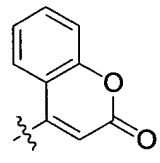


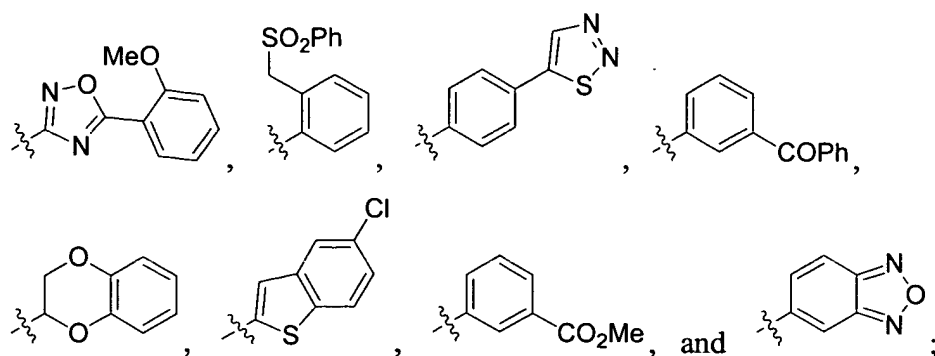
R<sup>2</sup> is R<sup>6</sup>, -CH<sub>2</sub>Ar<sup>1</sup>, -CHPh<sub>2</sub>, -CH<sub>2</sub>CO(4-FPh), -CH<sub>2</sub>CO(4-CF<sub>3</sub>Ph), or  
-CH<sub>2</sub>CONp where Np is naphthyl;

R<sup>3</sup> is C<sub>5-7</sub>cycloalkyl;

R<sup>4</sup> is hydrogen, Ar<sup>2</sup>, or Ar<sup>3</sup>;

Ar<sup>1</sup> is selected from the following group: phenyl, halophenyl,





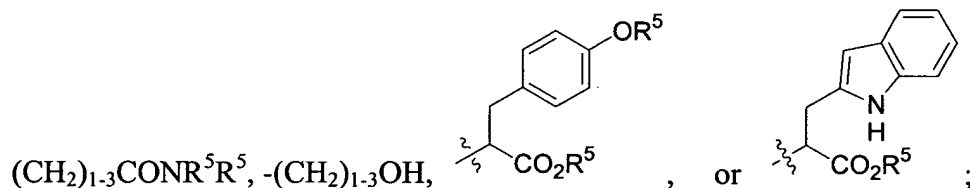
Ar<sup>2</sup> is phenyl, naphthyl, or biphenyl, optionally substituted with 1-3 substituents selected from the group comprising halogen, C<sub>1-6</sub> alkyl, hydroxyC<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, C<sub>1-6</sub>sulfoxy, C<sub>1-2</sub>perfluoroalkyl, hydroxy, formyl, C<sub>1-6</sub>alkylcarbonyl, cyano, nitro, C<sub>1-6</sub>alkylamido, CO<sub>2</sub>R<sup>5</sup>, CONR<sup>5</sup>R<sup>5</sup>, C<sub>1-6</sub>alkylsulfonamido, and dioxolane;

Ar<sup>3</sup> is thienyl, furanyl, pyrrolyl, benzothiophenyl, benzofuranyl, indolyl, quinoliny, or pyrimidinyl optionally substituted with 1-2 substituents selected from the group comprising C<sub>1-6</sub>alkyl, formyl, acetoxy, trifluoroacetoxy, and t-butoxycarbonyl;

R<sup>5</sup> is hydrogen or C<sub>1-6</sub>alkyl;

R<sup>6</sup> is halogen, methoxy, CO<sub>2</sub>R<sup>5</sup> or CONR<sup>7</sup>R<sup>8</sup>;

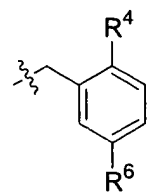
R<sup>7</sup> and R<sup>8</sup> are independently hydrogen, C<sub>1-6</sub>alkyl, -CH(Me)CO<sub>2</sub>R<sup>5</sup>, -(CH<sub>2</sub>)<sub>1-3</sub>CO<sub>2</sub>R<sup>5</sup>, -



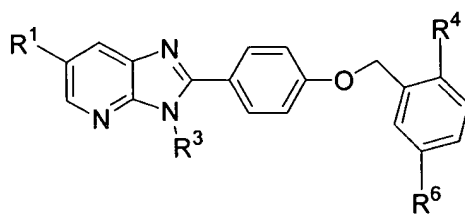
(CH<sub>2</sub>)<sub>1-3</sub>CONR<sup>5</sup>R<sup>5</sup>, -(CH<sub>2</sub>)<sub>1-3</sub>OH, or R<sup>7</sup> and R<sup>8</sup> taken together with the nitrogen to which they are attached form pyrrolidine, morpholine, piperidine, 4-hydroxypiperidine, piperazine, or 4-methylpiperazine;

or a pharmaceutically acceptable salt, solvate, or prodrug thereof.

2. (Original) A compound of claim 1 wherein R<sup>3</sup> is cyclohexyl.



3. (Original) A compound of claim 1 wherein R<sup>1</sup> is tetrazolyl and R<sup>2</sup> is
4. (Original) A compound of claim 3 wherein R<sup>4</sup> is Ar<sup>2</sup>.
5. (Original) A compound of claim 4 wherein R<sup>3</sup> is cyclohexyl.
6. (Original) A compound of claim 3 wherein R<sup>4</sup> is Ar<sup>3</sup>.
7. (Original) A compound of claim 6 wherein R<sup>3</sup> is cyclohexyl.
8. (Original) A compound of claim 3 wherein R<sup>4</sup> is hydrogen.
9. (Original) A compound of claim 8 wherein R<sup>3</sup> is cyclohexyl.
10. (Original) A compound of claim 1 wherein R<sup>2</sup> is -CH<sub>2</sub>Ar<sup>1</sup>.
11. (Original) A compound of claim 10 wherein R<sup>3</sup> is cyclohexyl.
12. (Original) A composition useful for treating hepatitis C comprising a therapeutic amount of a compound of claim 1 and a pharmaceutically acceptable carrier.
13. (Original) A method for treating hepatitis C comprising administering a therapeutically effective amount of a compound of claim 1 to a patient.
14. (New) A compound of Formula Ia



Ia

wherein:

R<sup>1</sup> is tetrazolyl or MeCONHSO<sub>2</sub>-;

R<sup>3</sup> is C<sub>5-7</sub>cycloalkyl;

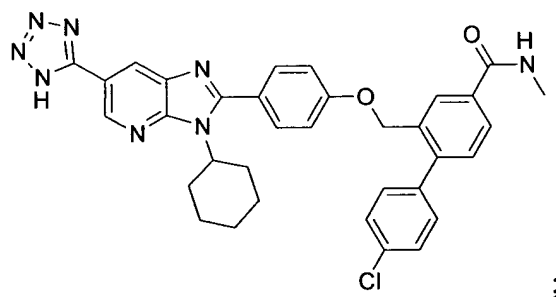
R<sup>4</sup> is phenyl substituted with halogen or cyano;

R<sup>6</sup> is methoxy or CONR<sup>7</sup>R<sup>8</sup>;

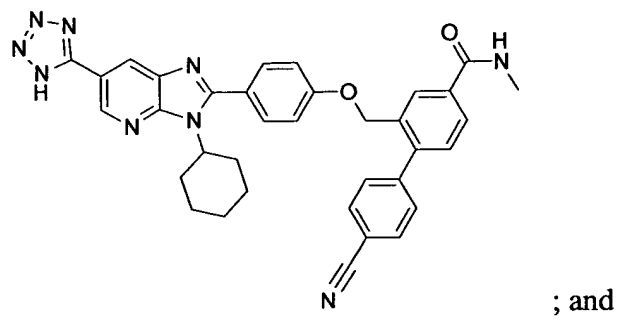
R<sup>7</sup> and R<sup>8</sup> are independently hydrogen or C<sub>1-6</sub>alkyl;

or a pharmaceutically acceptable salt, solvate, or prodrug thereof.

15. (New) A compound of claim 14 selected from the group consisting of;



;



; and

